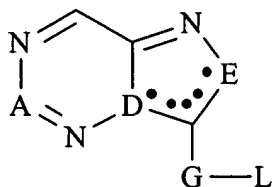


**IN THE CLAIMS:**

1. (Currently Amended) A method for inhibiting adenosine monophosphate deaminase (AMPDA) or adenosine deaminase (ADA) of plants which comprises applying compound of the formula (I), its tautomer, its salt or its water addition product,



(I)

where in formula (I)

A is a nitrogen atom or a group of the formula C-R, where R is as defined further below,

D is a carbon atom or a nitrogen atom,

E a) in the case that D is a nitrogen atom, is a nitrogen atom or a group of the formula C-R<sup>O</sup>, where R<sup>O</sup> is as defined further below, or

b) in the case that D is a carbon atom, is a group of the formula N- R<sup>O</sup>, -O-, -S-, -SO- or -SO<sub>2</sub>-,

the line of dots (•••••) from D via an adjacent ring carbon atom to E is a double bond

between the ring carbon atom and E if D is a nitrogen atom (case a), or

is a double bond between the ring carbon atom and D if D is a carbon atom (case

b),

R, R<sup>O</sup> independently of one another are each a hydrogen atom, amino, hydroxyl, mercapto, cyano, halogen, azido, nitro, SF<sub>5</sub>, unsubstituted or substituted

aminosulfonyl, acyl, acylamino, acyloxy, acylthio, mono- or di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, mono- or di(C<sub>3</sub>-C<sub>9</sub>)cycloalkylamino, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>2</sub>-C<sub>4</sub>)alkenylthio, (C<sub>2</sub>-C<sub>4</sub>)alkynylthio, (C<sub>3</sub>-C<sub>9</sub>)cycloalkylthio, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenylthio, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfinyl, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfonyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>2</sub>-C<sub>4</sub>)alkenyloxy, (C<sub>2</sub>-C<sub>4</sub>)alkynyloxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyloxy, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>2</sub>-C<sub>4</sub>)alkenyl, (C<sub>2</sub>-C<sub>4</sub>)alkynyl, (C<sub>3</sub>-C<sub>9</sub>) cycloalkyl, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyl, (C<sub>1</sub>-C<sub>4</sub>)alkylaminosulfonyl or di[(C<sub>1</sub>-C<sub>4</sub>)alkyl]aminosulfonyl, where each of the 23 last-mentioned radicals is unsubstituted or substituted in the hydrocarbon moiety by one or more radicals selected from the group consisting of halogen, hydroxyl, amino, nitro, formyl, carboxy, cyano, thiocyanato, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)haloalkylthio, mono(C<sub>1</sub>-C<sub>4</sub>)alkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>3</sub>-C<sub>9</sub>)cycloalkylamino, [(C<sub>1</sub>-C<sub>4</sub>)alkyl]carbonyl, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonyl, aminocarbonyl, mono(C<sub>1</sub>-C<sub>4</sub>)alkylaminocarbonyl and di(C<sub>1</sub>-C<sub>4</sub>)alkylaminocarbonyl,

- G is a divalent straight-chain saturated or unsaturated hydrocarbon bridge having 4 to 6 carbon atoms in the chain, in which one or more chain members, in each case independently of one another, can be replaced by O, S, NH, (C<sub>1</sub>-C<sub>4</sub>)alkyl-N or acyl-N or, in the unsaturated case, one or more CH groups can in each case be replaced by a nitrogen atom, where the bridge in question is unsubstituted or
- (a) substituted by one or more identical or different radicals selected from the group consisting of halogen, nitro, radicals of the formula R<sup>1</sup> which are

different from hydrogen, radicals of the formula  $R^2R^3C=$  and radicals of the formula  $L^*$ , where  $R^1$ ,  $R^2$ ,  $R^3$  and  $L^*$  are as defined further below,

- (b) carries two or four substituents, of which in each case two together with the linking bridge moiety form a carbocyclic or heterocyclic ring having 3 to 7 ring atoms, where in the case of a heterocycle the heteroatoms, preferably 1, 2 or 3 heteroatoms, are selected from the group consisting of N, O and S and where the ring in question may also have fused-on rings and is otherwise unsubstituted or substituted by one or more identical or different radicals selected from the group consisting of halogen, nitro, radicals of the formula  $R^1$  which are different from hydrogen, radicals of the formula  $L^*$  and oxo, where  $R^1$  and  $L^*$  are as defined further below,
- (c) is linked cyclically with L via a second direct bond or via a heteroatom selected from the group consisting of N, O and S,
- (d) has two or more substituents from the above groups (a) to (c) together,

L,  $L^*$  independently or one another are each  $OR^4$ ,  $SR^4$ , CN, tetrazolo,

$C(OR^5)(OR^6)OR^7$ ,  $-Z^1$ ,  $-O-Z^2$  or  $-NH-Z^2$ , where  $R^4$ ,  $R^5$ ,  $R^6$ ,  $R^7$ ,  $Z^1$  and  $Z^2$  are as defined further below and where L may be attached cyclically to the bridge G via a second direct bond or via a heteroatom selected from the group consisting of N, O and S,

$Z^1$  is a radical of the formula  $COOR^8$ ,  $CS-OR^8$ ,  $CO-SR^8$ ,  $CS-SR^8$ ,  $CO-NR^9-SO_2-R^8$ ,  $CO-NR^{10}R^{11}$ ,  $CS-NR^{10}R^{11}$ ,  $CO-R^{12}$ ,  $CS-R^{12}$ ,  $SO-R^{12}$ ,  $SO_2R^{12}$ ,  $SO_3R^8$ ,  $SO_2NR^{10}R^{11}$ ,  $SO_2NR^9COR^{12}$ ,  $SO_2NR^9COOR^{12}$ ,  $P(=O)(OR^{13})(OR^{14})$ ,

$P(=S)(OR^{13})(OR^{14})$ ,  $P(=O)(R^{15})(OR^{14})$ ,  $P(=O)(OR^{13})(NR^{10}R^{11})$ ,  $P(=O)(NR^{10}R^{11})-$   
 $(NR^{16}R^{17})$ ,  $P(=S)(OR^{13})(NR^{10}R^{11})$  or  $P(=S)(NR^{10}R^{11})(NR^{16}R^{17})$ ,

~~where the radical is formally formed by removing the hydroxyl group from the acid function~~

$Z^2$  is a radical of the formula  $COOR^8$ ,  $CS-OR^8$ ,  $CO-SR^8$ ,  $CS-SR^8$ ,  $CO-NR^9-SO_2R^8$ ,  
 $CO-NR^{10}R^{11}$ ,  $CS-NR^{10}R^{11}$ ,  $CO-R^{12}$ ,  $CS-R^{12}$ ,  $SO-R^{12}$ ,  $SO_2R^{12}$ ,  $SO_3R^8$ ,  
 $SO_2NR^{10}R^{11}$ ,  $SO_2NR^9COR^{12}$ ,  $SO_2NR^9COOR^{12}$ ,  $P(=O)(OR^{13})(OR^{14})$ ,  
 $P(=S)(OR^{13})(OR^{14})$ ,  $P(=O)(R^{15})(O^{14})$ ,  $P(=O)(OR^{13})(NR^{10}R^{11})$ ,  $P(=O)(R^{10}R^{11})-$   
 $(NR^{16}R^{17})$ ,  $P(=S)(OR^{13})(NR^{10}R^{11})$  or  $P(=S)(NR^{10}R^{11})(NR^{16}R^{17})$ ,

~~where the radical is formally formed by removing the hydroxyl group from the acid function,~~

$R^1$  to  $R^{17}$  independently of one another are each a hydrogen atom,  $(C_1-C_6)$ alkyl,  $(C_2-C_6)$ alkenyl,  $(C_2-C_6)$ alkynyl,  $(C_3-C_9)$ cycloalkyl,  $(C_5-C_9)$ cycloalkenyl, phenyl or heterocyclyl, where each of the last-mentioned carbon-containing radicals is unsubstituted or substituted by one or more radicals selected from the group consisting of amino, hydroxyl, mercapto, cyano, halogen, azido, nitro,  $SF_5$ , aminosulfonyl, acyl, acylamino, acyloxy, acylthio,  $[(C_1-C_4)$ alkoxy]carbonyl, mono $(C_1-C_4)$ alkylamino, mono $(C_3-C_9)$ cycloalkylamino, di $(C_1-C_4)$ alkylamino,  $(C_1-C_4)$ alkylthio,  $(C_2-C_4)$ alkenylthio,  $(C_2-C_4)$ alkynylthio,  $(C_3-C_9)$ cycloalkylthio,  $(C_5-C_9)$ cycloalkenylthio,  $(C_1-C_4)$ alkylsulfinyl,  $(C_1-C_4)$ alkylsulfonyl,  $(C_1-C_4)$ alkoxy,  $(C_2-C_4)$ alkenyloxy,  $(C_2-C_4)$ alkynyloxy,  $(C_3-C_9)$ cycloalkoxy,  $(C_5-C_9)$ cycloalkenyloxy,  $(C_3-C_9)$ cycloalkyl,  $(C_5-C_9)$ cycloalkenyl, phenyl, substituted phenyl, heteroaryl, substituted heteroaryl and, in the case of cyclic radicals, also

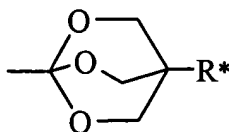
by (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>2</sub>-C<sub>4</sub>)alkenyl, (C<sub>2</sub>-C<sub>4</sub>)alkynyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>2</sub>-C<sub>4</sub>)haloalkenyl, (C<sub>2</sub>-C<sub>4</sub>)haloalkynyl, (C<sub>1</sub>-C<sub>4</sub>)hydroxyalkyl and (C<sub>1</sub>-C<sub>4</sub>)alkoxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, where heterocyclyl in the definition of R<sup>1</sup> to R<sup>17</sup> is a radical of is a heterocyclic saturated, unsaturated or heteroaromatic ring having 3 to 6 ring atoms and 1 to 3 heteroatoms selected from the group consisting of N, O and S

where heteroaryl in the definition of R<sup>1</sup> to R<sup>17</sup> is a radical of is a heteroaromatic ring having 5 to 6 ring atoms and 1 to 3 heteroatoms selected from the group consisting of N, O and S and where the substituents for substituted phenyl or substituted heteroaryl are one or more radicals selected from the group consisting of halogen, nitro, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)hydroxyalkyl and (C<sub>1</sub>-C<sub>4</sub>)alkoxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, or

R<sup>2</sup>, R<sup>3</sup> together with the carbon atom of the group R<sup>2</sup>R<sup>3</sup>C= are a non-aromatic carbocyclic ring or a heterocyclic ring having 3 to 9 ring atoms and 1 to 4 heteroring atoms selected from the group consisting of N, O and S, which ring is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, nitro, hydroxyl, oxo, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy and (C<sub>1</sub>-C<sub>4</sub>)alkylthio, or

R<sup>5</sup>, R<sup>6</sup> together with the carbon atom and the adjacent oxygen atoms of the group C(OR<sup>5</sup>)(OR<sup>6</sup>)(OR<sup>7</sup>) are a saturated or unsaturated non-aromatic heterocyclic ring having 3 to 9 ring atoms and 1 to 4 heteroring atoms selected from the group consisting of N, O, P, and S, which ring is unsubstituted or substituted by one or

more radicals selected from the group consisting of halogen, nitro, hydroxyl, oxo, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy and (C<sub>1</sub>-C<sub>4</sub>)alkylthio, or the group C(OR<sup>5</sup>)(OR<sup>6</sup>)(OR<sup>7</sup>) together is a bicyclic radical of the formula



in which

R\* is (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio or phenyl which is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, nitro, hydroxyl, oxo, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy and (C<sub>1</sub>-C<sub>4</sub>)alkylthio,

to the enzyme AMPA AMPDA of plants or enzyme ADA of plants.

2. (Previously Presented) The method as claimed in claim 1, wherein

A is a nitrogen atom or

a group of the formula C-R in which

R is a hydrogen atom, amino, hydroxyl, mercapto, cyano, halogen, azido, nitro, SF<sub>5</sub>, aminosulfonyl, (C<sub>1</sub>-C<sub>5</sub>)alkanoylamino, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonylamino, (C<sub>1</sub>-C<sub>5</sub>)alkanoyl, [(C<sub>1</sub>-C<sub>4</sub>)-alkoxy]carbonyl, (C<sub>1</sub>-C<sub>5</sub>)alkanoyloxy, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonyloxy, mono-(C<sub>1</sub>-C<sub>4</sub>)alkylamino, mono(C<sub>3</sub>-C<sub>6</sub>)cycloalkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfinyl, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfonyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>3</sub>-C<sub>4</sub>)alkenyloxy, (C<sub>3</sub>-C<sub>4</sub>)alkynyloxy, (C<sub>3</sub>-C<sub>6</sub>)cycloalkoxy, (C<sub>3</sub>-

C<sub>6</sub>)cycloalkenyloxy, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>2</sub>-C<sub>4</sub>)alkenyl, (C<sub>2</sub>-C<sub>4</sub>)alkynyl, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, (C<sub>5</sub>-C<sub>6</sub>)cycloalkenyl, (C<sub>1</sub>-C<sub>4</sub>)alkylaminosulfonyl or di[(C<sub>1</sub>-C<sub>4</sub>)alkyl]aminosulfonyl, where each of the 24 last-mentioned radicals is unsubstituted or substituted in the hydrocarbon moiety by one or more radicals selected from the group consisting of halogen, hydroxyl, amino, nitro, formyl, carboxyl, cyano, thiocyanato, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>3</sub>-C<sub>6</sub>)cycloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)haloalkylthio, mono(C<sub>1</sub>-C<sub>4</sub>)alkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, (C<sub>3</sub>-C<sub>6</sub>)cycloalkylamino, [(C<sub>1</sub>-C<sub>4</sub>)alkyl]carbonyl, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonyl, aminocarbonyl, mono(C<sub>1</sub>-C<sub>4</sub>)alkylaminocarbonyl and di(C<sub>1</sub>-C<sub>4</sub>)alkylaminocarbonyl.

3. (Currently Amended)      The method as claimed in claim 1, wherein in the compound of formula (I)

G      is a divalent straight-chain saturated or unsaturated hydrocarbon bridge having 4 to 6 carbon atoms in the chain, in which one or more CH<sub>2</sub> groups, in each case independently of one another, are replaced by O or S, where the bridge in question is unsubstituted or

- (a)      substituted by one or more halogen atoms and additionally or alternatively by one or more identical or different radicals selected from the group consisting of nitro, radicals of the formula R<sup>1</sup> which are different from hydrogen, radicals of the formula R<sup>2</sup>R<sup>3</sup>C= and radicals of the formula L\*, where R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and L\* are as defined above,
- (b)      carries two or four substituents, in each case two of which together with the linking bridge moiety form a carbocyclic ring having 3 to 6 carbon

atoms or a heterocyclic saturated or partially unsaturated ring having 3 to 6 ring atoms or a heteroaromatic ring having 5 or 6 ring atoms, where in the case of a heterocycle, the 1, 2 or 3 heteroatoms are selected from the group consisting of N, O and S and where the ring in question may also have a fused-on carbocyclic ring having 4 to 6 ring atoms or a fused-on heterocyclic ring having 4 to 6 ring atoms and 1, 2 or 3 heteroatoms selected from the group consisting of N, O and S, the ring being otherwise unsubstituted or substituted by one or more halogen atoms and additionally or alternatively by one or more identical or different radicals selected from the group consisting of nitro, radicals or the formula  $R^1$  which are different from hydrogen, radicals of the formula  $L^*$  and oxo, where  $R^1$  and  $L^*$  are as defined above,

- (c) has substituents from the ~~above~~ as defined in the compound of formula I groups (a) and (b) together.

4. (Cancelled)

5. (Currently Amended) The method as claimed in claim 1, wherein

G is a divalent straight-chain saturated or unsaturated hydrocarbon bridge having ~~to 8~~ 4 to 6 carbon atoms in the chain in which one or more  $CH_2$  groups, in each case independently of one another, are replaced by O or S,

or is a bridge of the formula  $-W^1$ -cycle- $W^2$ -, in which

$W^1$ ,  $W^2$  independently of one another are a direct bond,  $CH_2$ ,  $CH_2CH_2$ ,  $OCH_2$ ,  $SCH_2$ ,  $CH_2CH_2CH_2$ ,  $CH_2OCH_2$ ,  $CH_2SCH_2$ ,  $OCH_2CH_2$  or  $SCH_2CH_2$  and "cycle" is 1,4-



cyclohexylene, 1,2-phenylene, 1,3-phenylene, 1,4-phenylene, 1,2-naphthylene, 1,3-naphthylene, 1,4-naphthylene, 1,2-tetrahydronaphthylene, 1,3-tetrahydronaphthylene, 1,4-tetrahydronaphthylene, 1,2-cyclopentylene, 1,3-cyclopentylene, 1,2-cyclohexylene, 1,3-cyclohexylene, 1,4-cyclohexylene, tetrahydrofuran-2,5-diyl (oxolane), tetrahydrothiophene-2,5-diyl, 2,5-dihydrofuran-2,5-diyl or 2,5-dihydrothiophene-2,5-diyl,

where the bridge in question is unsubstituted or

substituted by one or more halogen atoms and additionally or alternatively by one or more identical or different radicals selected from the group consisting of radicals of the formula  $R^1$  which are different from hydrogen, radicals of the formula  $R^2R^3C=$  and radicals of the formula  $L^*$ , where  $R^1$ ,  $R^2$ ,  $R^3$  and  $L^*$  are as defined above or further below, or

is additionally or alternatively attached cyclically to L via a second direct bond or via a heteroatom selected from the group consisting of N, O and S, and

$R^1$  to  $R^{17}$  independently of one another are each a hydrogen atom,  $(C_1-C_4)$ alkyl,  $(C_2-C_4)$ alkenyl,  $(C_2-C_4)$ alkynyl,  $(C_3-C_6)$ cycloalkyl,  $(C_5-C_6)$ cycloalkenyl, phenyl or heterocyclyl, where each of the last-mentioned radicals is unsubstituted or substituted by one or more radicals selected from the group consisting of amino, hydroxyl, mercapto, cyano, halogen, azido, nitro,  $SF_5$ , aminosulfonyl,  $(C_1-C_4)$ alkanoyl,  $(C_1-C_4)$ alkanoylamino, benzoylamino,  $(C_1-C_4)$ alkanoyloxy,  $(C_1-C_4)$ alkanoylthio,  $[C_1-C_4]$ alkoxy]carbonyl, mono( $C_1-C_4$ )alkylamino, di( $C_1-C_4$ )alkylamino,  $(C_1-C_4)$ alkylthio,  $(C_3-C_4)$ alkenylthio,  $(C_3-C_4)$ alkynylthio,  $(C_1-C_4)$ alkylsulfinyl,  $(C_1-C_4)$ alkylsulfonyl,  $(C_1-C_4)$ alkoxy,  $(C_3-C_4)$ alkenyloxy,  $(C_3-$

(C<sub>4</sub>)alkynyloxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, phenyl, substituted phenyl, heteroaryl, substituted heteroaryl and, in the case of cyclic radicals, also by (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>2</sub>-C<sub>4</sub>)alkenyl, (C<sub>2</sub>-C<sub>4</sub>)alkynyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>2</sub>-C<sub>4</sub>)haloalkenyl, (C<sub>2</sub>-C<sub>4</sub>)halo-alkynyl, (C<sub>1</sub>-C<sub>4</sub>)hydroxyalkyl and (C<sub>1</sub>-C<sub>4</sub>)alkoxyl(C<sub>1</sub>-C<sub>4</sub>)alkyl, where heterocyclyl is a heterocyclic saturated or unsaturated ring having 3 to 6 ring atoms or a heteroaromatic ring having 5 or 6 ring atoms and in each case 1 to 3 heteroatoms selected from the group consisting of N, O and S and where heteroaryl is a heteroaromatic ring having 5 to 6 ring atoms and 1 to 3 heteroatoms selected from the group consisting of N, O and S and where the substituents for substituted phenyl or substituted heteroaryl are one or more substituents selected from the group consisting of halogen, nitro, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)hydroxyalkyl and (C<sub>1</sub>-C<sub>4</sub>)alkoxy(C<sub>1</sub>-C<sub>4</sub>)alkyl.

6. (Previously Presented) The method as claimed in claim 1, wherein

L is hydroxyl, carboxyl, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonyl, CONH<sub>2</sub>, [(C<sub>1</sub>-C<sub>4</sub>)alkylamino]carbonyl, [(C<sub>1</sub>-C<sub>4</sub>)alkylsulfonylamino]carbonyl, [(C<sub>1</sub>-C<sub>4</sub>)haloalkylsulfonylamino]carbonyl, [cyano(C<sub>1</sub>-C<sub>4</sub>)alkylsulfonylamino]carbonyl, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfonylamino, (C<sub>1</sub>-C<sub>4</sub>)haloalkylsulfonylamino, cyano-(C<sub>1</sub>-C<sub>4</sub>)alkylsulfonylamino, (C<sub>1</sub>-C<sub>5</sub>)alkanoyloxy, benzyloxy, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonyloxy, [(C<sub>1</sub>-C<sub>4</sub>)alkylamino]carbonyloxy, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)hydroxyalkoxy, SO<sub>2</sub>NHCONH<sub>2</sub>, (C<sub>1</sub>-C<sub>5</sub>)alkanoylamino-sulfonyl, [(C<sub>1</sub>-C<sub>4</sub>)haloalkyl]carbonylamino-sulfonyl, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]-

carbonylaminosulfonyl,  $[\text{C}_1\text{-C}_5]\text{haloalkoxy}$ carbonylaminosulfonyl,  $\text{SO}_2\text{NH}_2$ ,  
 $\text{di}[(\text{C}_1\text{-C}_4)\text{alkyl}]\text{aminosulfonyl}$ ,  $\text{P}(=\text{O})(\text{OH})_2$ ,  $\text{P}(=\text{S})(\text{OH})_2$ ,  $\text{P}(=\text{O})(\text{OR}')_2$  or  
 $\text{P}(=\text{O})(\text{OH})(\text{OR}')$ , where in the two last mentioned formulae  $\text{R}'$ , in each case  
independently of any other radicals  $\text{R}'$ , is  $(\text{C}_1\text{-C}_4)\text{alkyl}$ ,  $(\text{C}_1\text{-C}_4)\text{haloalkyl}$ ,  $(\text{C}_1\text{-C}_4)\text{hydroxyalkyl}$ ,  $(\text{C}_1\text{-C}_4)\text{alkanoyl}(\text{C}_1\text{-C}_4)\text{alkyl}$ ,  $(\text{C}_1\text{-C}_4)\text{alkanoyloxy}(\text{C}_1\text{-C}_4)\text{alkyl}$   
or phenyl.

7. (Currently Amended) A compound of the formula (I),



where in formula (I)

A is a nitrogen atom or a group of the formula C-R, where R is as defined further below,

D is a carbon atom or a nitrogen atom,

E a) in the case that D is a nitrogen atom, is a nitrogen atom or a group of the formula C-R<sup>O</sup>, where R<sup>O</sup> is as defined further below, or

b) in the case that D is a carbon atom, is a group of the formula N- R<sup>O</sup>, -O-, -S-, -SO- or -SO<sub>2</sub>-,

the line of dots (•••••) from D via an adjacent ring carbon atom to E is a double bond between the ring carbon atom and E if D is a nitrogen atom (case a), or

is a double bond between the ring carbon atom and D if D is a carbon atom (case b),

R, R<sup>O</sup> independently of one another are each a hydrogen atom, amino, hydroxyl, mercapto, cyano, halogen, azido, nitro, SF<sub>5</sub>, unsubstituted or substituted aminosulfonyl, acyl, acylamino, acyloxy, acylthio, mono- or di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, mono- or di(C<sub>3</sub>-C<sub>9</sub>)cycloalkylamino, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>2</sub>-C<sub>4</sub>)alkenylthio, C<sub>2</sub>-C<sub>4</sub>alkynylthio, (C<sub>3</sub>-C<sub>9</sub>)cycloalkylthio, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenylthio, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfinyl, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfonyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>2</sub>-C<sub>4</sub>)alkenyloxy, (C<sub>2</sub>-C<sub>4</sub>)alkynyloxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyloxy, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>2</sub>-C<sub>4</sub>)alkenyl, (C<sub>2</sub>-C<sub>4</sub>)alkynyl, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyl, (C<sub>1</sub>-C<sub>4</sub>)alkylaminosulfonyl or di[(C<sub>1</sub>-C<sub>4</sub>)alkyl]aminosulfonyl, where each of the 23 last-mentioned radicals is unsubstituted or substituted in the hydrocarbon moiety by one or more radicals selected from the group consisting of halogen, hydroxyl, amino, nitro, formyl, carboxy, cyano, thiocyanato, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)haloalkylthio, mono(C<sub>1</sub>-C<sub>4</sub>)alkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>3</sub>-C<sub>9</sub>)cycloalkylamino, [(C<sub>1</sub>-C<sub>4</sub>)alkyl]carbonyl, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonyl, aminocarbonyl, mono(C<sub>1</sub>-C<sub>4</sub>)alkylaminocarbonyl and di(C<sub>1</sub>-C<sub>4</sub>)alkylaminocarbonyl,

G is a divalent straight-chain saturated or unsaturated hydrocarbon bridge having 4 to 6 carbon atoms in the chain, in which one or more chain members, in each case independently of one another, can be replaced by O, S, NH, (C<sub>1</sub>-C<sub>4</sub>)alkyl-N or

acyl-N or, in the unsaturated case, one or more CH groups can in each case be replaced by a nitrogen atom, where the bridge in question is unsubstituted or

(a) substituted by one or more identical or different radicals selected from the group consisting of halogen, nitro, radicals of the formula  $R^1$  which are different from hydrogen, radicals of the formula  $R^2R^3C=$  and radicals of the formula  $L^*$ , where  $R^1$ ,  $R^2$ ,  $R^3$  and  $L^*$  are as defined further below,

(b) carries two or four substituents, of which in each case two together with the linking bridge moiety form a carbocyclic or heterocyclic ring having 3 to 7 ring atoms, where in the case of a heterocycle the heteroatoms, preferably 1, 2 or 3 heteroatoms, are selected from the group consisting of N, O and S and where the ring in question may also have fused-on rings and is otherwise unsubstituted or substituted by one or more identical or different radicals selected from the group consisting of halogen, nitro, radicals of the formula  $R^1$  which are different from hydrogen, radicals of the formula  $L^*$  and oxo, where  $R^1$  and  $L^*$  are as defined further below,

(c) is linked cyclically with L via a second direct bond or via a heteroatom selected from the group consisting of N, O and S,

(d) has two or more substituents from the above groups (a) to (c) together,

L,  $L^*$  independently or one another are each  $OR^4$ ,  $SR^4$ , CN, tetrazolo,

$C(OR^5)(OR^6)OR^7$ ,  $-Z^1$ ,  $-O-Z^2$  or  $-NH-Z^2$ , where  $R^4$ ,  $R^5$ ,  $R^6$ ,  $R^7$ ,  $Z^1$  and  $Z^2$  are as defined further below and where L may be attached cyclically to the bridge G via a second direct bond or via a heteroatom selected from the group consisting of, O and S,

Z<sup>1</sup> is a radical of the formula COOR<sup>8</sup>, CS-OR<sup>8</sup>, CO-SR<sup>8</sup>, CS-SR<sup>8</sup>, CO-NR<sup>9</sup>-SO<sub>2</sub>-R<sup>8</sup>, CO-NR<sup>10</sup>R<sup>11</sup>, CS-NR<sup>10</sup>R<sup>11</sup>, CO-R<sup>12</sup>, CS-R<sup>12</sup>, SO-R<sup>12</sup>, SO<sub>2</sub>R<sup>12</sup>, SO<sub>3</sub>R<sup>8</sup>, SO<sub>2</sub>NR<sup>10</sup>R<sup>11</sup>, SO<sub>2</sub>NR<sup>9</sup>COR<sup>12</sup>, SO<sub>2</sub>NR<sup>9</sup>COOR<sup>12</sup>, P(=O)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=S)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=O)(R<sup>15</sup>)(OR<sup>14</sup>), P(=O)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>), P(=O)(NR<sup>10</sup>R<sup>11</sup>)-(NR<sup>16</sup>R<sup>17</sup>), P(=S)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>) or P(=S)(NR<sup>10</sup>R<sup>11</sup>)(NR<sup>16</sup>R<sup>17</sup>).

Z<sup>2</sup> is a radical of the formula COOR<sup>8</sup>, CS-OR<sup>8</sup>, CO-SR<sup>8</sup>, CS-SR<sup>8</sup>, CO-NR<sup>9</sup>-SO<sub>2</sub>-R<sup>8</sup>, CO-NR<sup>10</sup>R<sup>11</sup>, CS-NR<sup>10</sup>R<sup>11</sup>, CO-R<sup>12</sup>, CS-R<sup>12</sup>, SO-R<sup>12</sup>, SO<sub>2</sub>R<sup>12</sup>, SO<sub>3</sub>R<sup>8</sup>, SO<sub>2</sub>NR<sup>10</sup>R<sup>11</sup>, SO<sub>2</sub>NR<sup>9</sup>COR<sup>12</sup>, SO<sub>2</sub>NR<sup>9</sup>COOR<sup>12</sup>, P(=O)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=S)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=O)(R<sup>15</sup>)(OR<sup>14</sup>), P(=O)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>), P(=O)(NR<sup>10</sup>R<sup>11</sup>)-(NR<sup>16</sup>R<sup>17</sup>), P(=S)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>) or P(=S)(NR<sup>10</sup>R<sup>11</sup>)(NR<sup>16</sup>R<sup>17</sup>).

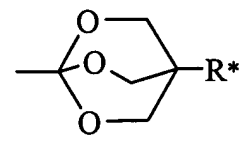
R<sup>1</sup> to R<sup>17</sup> independently of one another are each a hydrogen atom, (C<sub>1</sub>-C<sub>6</sub>)alkyl, (C<sub>2</sub>-C<sub>6</sub>)alkenyl, (C<sub>2</sub>-C<sub>6</sub>)alkynyl, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyl, phenyl or heterocyclyl, where each of the last-mentioned carbon-containing radicals is unsubstituted or substituted by one or more radicals selected from the group consisting of amino, hydroxyl, mercapto, cyano, halogen, azido, nitro, SF<sub>5</sub>, aminosulfonyl, acyl, acylamino, acyloxy, acylthio, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonyl, mono(C<sub>1</sub>-C<sub>4</sub>)alkylamino, mono(C<sub>3</sub>-C<sub>9</sub>)cycloalkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>2</sub>-C<sub>4</sub>)alkenylthio, (C<sub>2</sub>-C<sub>4</sub>)alkynylthio, (C<sub>3</sub>-C<sub>9</sub>)cycloalkylthio, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenylthio, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfinyl, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfonyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>2</sub>-C<sub>4</sub>)alkenyloxy, (C<sub>2</sub>-C<sub>4</sub>)alkynyloxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyloxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyl, phenyl, substituted phenyl, heteroaryl, substituted heteroaryl and, in the case of cyclic radicals, also by (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>2</sub>-C<sub>4</sub>)alkenyl, (C<sub>2</sub>-C<sub>4</sub>)alkynyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>2</sub>-

(C<sub>4</sub>)haloalkenyl, (C<sub>2</sub>-C<sub>4</sub>)haloalkynyl, (C<sub>1</sub>-C<sub>4</sub>)hydroxyalkyl and (C<sub>1</sub>-C<sub>4</sub>)alkoxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, where heterocyclyl in the definition of R<sup>1</sup> to R<sup>17</sup> is a radical of a heterocyclic saturated, unsaturated or heteroaromatic ring having 3 to 6 ring atoms and 1 to 3 heteroatoms selected from the group consisting of N, O and S where heteroaryl in the definition of R<sup>1</sup> to R<sup>17</sup> is a radical of a heteroaromatic ring having 5 to 6 ring atoms and 1 to 3 heteroatoms selected from the group consisting of N, O and S and where the substituents for substituted phenyl or substituted heteroaryl are one or more radicals selected from the group consisting of halogen, nitro, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)hydroxyalkyl and (C<sub>1</sub>-C<sub>4</sub>)alkoxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, or

R<sup>2</sup>, R<sup>3</sup> together with the carbon atom of the group R<sup>2</sup>R<sup>3</sup>C= are a non-aromatic carbocyclic ring or a heterocyclic ring having 3 to 9 ring atoms and 1 to 4 heteroring atoms selected from the group consisting of N, O and S, which ring is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, nitro, hydroxyl, oxo, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy and (C<sub>1</sub>-C<sub>4</sub>)alkylthio, or

R<sup>5</sup>, R<sup>6</sup> together with the carbon atom and the adjacent oxygen atoms of the group C(OR<sup>5</sup>)(OR<sup>6</sup>)(OR<sup>7</sup>) are a saturated or unsaturated non-aromatic heterocyclic ring having 3 to 9 ring atoms and 1 to 4 heteroring atoms selected from the group consisting of N, O, P, and S, which ring is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, nitro, hydroxyl, oxo,

(C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy and (C<sub>1</sub>-C<sub>4</sub>)alkylthio, or the group C(OR<sup>5</sup>)(OR<sup>6</sup>)(OR<sup>7</sup>) together is a bicyclic radical of the formula



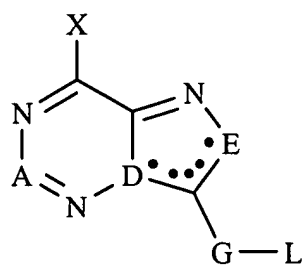
in which

R\* is (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio or phenyl which is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, nitro, hydroxyl, oxo, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy and (C<sub>1</sub>-C<sub>4</sub>)alkylthio,

its tautomers, its salts or its water addition product as defined in claim 1, except for the compound of the formula (I) in which A = CH, D = C, E = NH and G-L = β-D-ribofuranosyl.

8. (Currently Amended) A process for preparing a compound of the formula (I) or a salt thereof as claimed in claim 7, which comprises
- a) reducing a compound of the formula (II)

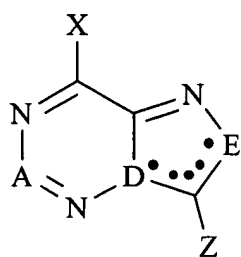




(II)

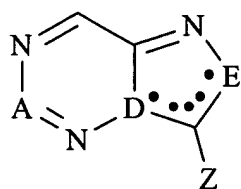
in which X is a leaving group to the compound of the formula (I) or

b) reducing a compound of the formula (III)



(III)

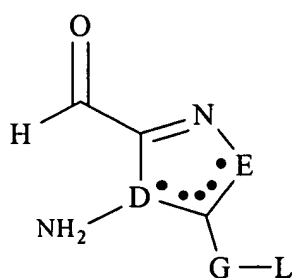
in which X is a leaving group and Z is a precursor of the radical G-L to the compound of the formula (III')



(III')

in which Z is as defined in formula (III), and then modifying the compound (III) at the group Z such that the compound (I) is obtained,

- c) modifying a compound of the formula (III') in which Z is a precursor of the radical G-L at the group Z such that the compound (I) is obtained, or
- d) if A is a group of the formula C-R, cyclizing a compound of the formula (III'')



(III'')

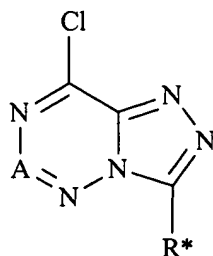
with a compound of the formula (III''')



in which A is a group C-R to give the compound of the formula (I),

where the symbols A, D, E, G, L and R in the formulae (II), (III) (III'), (III'') and (III''') are as defined in said compound of formula (I), unless specifically defined otherwise.

9. (Currently Amended) A process for preparing a compound of the formula (V)



(V)

in which  $R^* = Z$  or G-L, Z is a precursor of the radical G-L

and A, G and L are defined further below

A is a nitrogen atom or a group of the formula C-R, where R is as defined further below,

R, is a hydrogen atom, amino, hydroxyl, mercapto, cyano, halogen, azido, nitro, SF<sub>5</sub>, unsubstituted or substituted aminosulfonyl, acyl, acylamino, acyloxy, acylthio, mono- or di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, mono- or di(C<sub>3</sub>-C<sub>9</sub>)cycloalkylamino, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>2</sub>-C<sub>4</sub>)alkenylthio, (C<sub>2</sub>-C<sub>4</sub>)alkynylthio, (C<sub>3</sub>-C<sub>9</sub>)cycloalkylthio, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenylthio, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfinyl, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfonyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>2</sub>-C<sub>4</sub>)alkenyloxy, (C<sub>2</sub>-C<sub>4</sub>)alkynyloxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyloxy, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>2</sub>-C<sub>4</sub>)alkenyl, (C<sub>2</sub>-C<sub>4</sub>)alkynyl, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyl, (C<sub>1</sub>-C<sub>4</sub>)alkylaminosulfonyl or di[(C<sub>1</sub>-C<sub>4</sub>)alkyl]aminosulfonyl, where each of the 23 last-mentioned radicals is unsubstituted or substituted in the hydrocarbon moiety by one or more radicals selected from the group consisting of halogen, hydroxyl, amino, nitro, formyl, carboxy, cyano, thiocyanato, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)haloalkylthio, mono(C<sub>1</sub>-C<sub>4</sub>)alkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>3</sub>-C<sub>9</sub>)cycloalkylamino, [(C<sub>1</sub>-

(C<sub>4</sub>)alkyl]carbonyl, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonyl, aminocarbonyl, mono(C<sub>1</sub>-C<sub>4</sub>)alkylaminocarbonyl and di(C<sub>1</sub>-C<sub>4</sub>)alkylaminocarbonyl,

- G is a divalent straight-chain saturated or unsaturated hydrocarbon bridge having 4 to 6 carbon atoms in the chain, in which one or more chain members, in each case independently of one another, can be replaced by O, S, NH, (C<sub>1</sub>-C<sub>4</sub>)alkyl-N or acyl-N or, in the unsaturated case, one or more CH groups can in each case be replaced by a nitrogen atom, where the bridge in question is unsubstituted or
- (a) substituted by one or more identical or different radicals selected from the group consisting of halogen, nitro, radicals of the formula R<sup>1</sup> which are different from hydrogen, radicals of the formula R<sup>2</sup>R<sup>3</sup>C= and radicals of the formula L\*, where R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and L\* are as defined further below,
- (b) carries two or four substituents, of which in each case two together with the linking bridge moiety form a carbocyclic or heterocyclic ring having 3 to 7 ring atoms, where in the case of a heterocycle the heteroatoms, preferably 1, 2 or 3 heteroatoms, are selected from the group consisting of N, O and S and where the ring in question may also have fused-on rings and is otherwise unsubstituted or substituted by one or more identical or different radicals selected from the group consisting of halogen, nitro, radicals of the formula R<sup>1</sup> which are different from hydrogen, radicals of the formula L\* and oxo, where R<sup>1</sup> and L\* are as defined further below,
- (c) is linked cyclically with L via a second direct bond or via a heteroatom selected from the group consisting of O and S,
- (d) has two or more substituents from the above groups (a) to (c) together,

L, L\* independently or one another are each OR<sup>4</sup>, SR<sup>4</sup>, CN, tetrazolo,

C(OR<sup>5</sup>)(OR<sup>6</sup>)OR<sup>7</sup>), -Z<sup>1</sup>, -O-Z<sup>2</sup> or -NH-Z<sup>2</sup>, where R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, Z<sup>1</sup> and Z<sup>2</sup> are as defined further below and where L may be attached cyclically to the bridge G via a second direct bond or via a heteroatom selected from the group consisting of N, O and S,

Z<sup>1</sup> is a radical of the formula COOR<sup>8</sup>, CS-OR<sup>8</sup>, CO-SR<sup>8</sup>, CS-SR<sup>8</sup>, CO-NR<sup>9</sup>-SO<sub>2</sub>-R<sup>8</sup>, CO-NR<sup>10</sup>R<sup>11</sup>, CS-NR<sup>10</sup>R<sup>11</sup>, CO-R<sup>12</sup>, CS-R<sup>12</sup>, SO-R<sup>12</sup>, SO<sub>2</sub>R<sup>12</sup>, SO<sub>3</sub>R<sup>8</sup>, SO<sub>2</sub>NR<sup>10</sup>R<sup>11</sup>, SO<sub>2</sub>NR<sup>9</sup>COR<sup>12</sup>, SO<sub>2</sub>NR<sup>9</sup>COOR<sup>12</sup>, P(=O)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=S)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=O)(R<sup>15</sup>)(OR<sup>14</sup>), P(=O)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>), P(=O)(NR<sup>10</sup>R<sup>11</sup>)-(NR<sup>16</sup>R<sup>17</sup>), P(=S)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>) or P(=S)(NR<sup>10</sup>R<sup>11</sup>)(NR<sup>16</sup>R<sup>17</sup>),

Z<sup>2</sup> is a radical of the formula COOR<sup>8</sup>, CS-OR<sup>8</sup>, CO-SR<sup>8</sup>, CS-SR<sup>8</sup>, CO-NR<sup>9</sup>-SO<sub>2</sub>-R<sup>8</sup>, CO-NR<sup>10</sup>R<sup>11</sup>, CS-NR<sup>10</sup>R<sup>11</sup>, CO-R<sup>12</sup>, CS-R<sup>12</sup>, SO-R<sup>12</sup>, SO<sub>2</sub>R<sup>12</sup>, SO<sub>3</sub>R<sup>8</sup>, SO<sub>2</sub>NR<sup>10</sup>R<sup>11</sup>, SO<sub>2</sub>NR<sup>9</sup>COR<sup>12</sup>, SO<sub>2</sub>NR<sup>9</sup>COOR<sup>12</sup>, P(=O)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=S)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=O)(R<sup>15</sup>)(OR<sup>14</sup>), P(=O)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>), P(=O)(NR<sup>10</sup>R<sup>11</sup>)-(NR<sup>16</sup>R<sup>17</sup>), P(=S)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>) or P(=S)(NR<sup>10</sup>R<sup>11</sup>)(NR<sup>16</sup>R<sup>17</sup>),

R<sup>1</sup> to R<sup>17</sup> independently of one another are each a hydrogen atom, (C<sub>1</sub>-C<sub>6</sub>)alkyl, (C<sub>2</sub>-C<sub>6</sub>)alkenyl, (C<sub>2</sub>-C<sub>6</sub>)alkynyl, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyl, phenyl or heterocyclyl, where each of the last-mentioned carbon-containing radicals is unsubstituted or substituted by one or more radicals selected from the group consisting of amino, hydroxyl, mercapto, cyano, halogen, azido, nitro, SF<sub>5</sub>, aminosulfonyl, acyl, acylamino, acyloxy, acylthio, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonyl, mono(C<sub>1</sub>-C<sub>4</sub>)alkylamino, mono(C<sub>3</sub>-C<sub>9</sub>)cycloalkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>2</sub>-C<sub>4</sub>)alkenylthio, (C<sub>2</sub>-C<sub>4</sub>)alkynylthio, (C<sub>3</sub>-C<sub>9</sub>)cycloalkylthio,

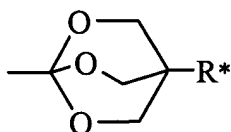
(C<sub>5</sub>-C<sub>9</sub>)cycloalkenylthio, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfinyl, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfonyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>2</sub>-C<sub>4</sub>)alkenyloxy, (C<sub>2</sub>-C<sub>4</sub>)alkynyloxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyloxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyl, phenyl, substituted phenyl, heteroaryl, substituted heteroaryl and, in the case of cyclic radicals, also by (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>2</sub>-C<sub>4</sub>)alkenyl, (C<sub>2</sub>-C<sub>4</sub>)alkynyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>2</sub>-C<sub>4</sub>)haloalkenyl, (C<sub>2</sub>-C<sub>4</sub>)haloalkynyl, (C<sub>1</sub>-C<sub>4</sub>)hydroxyalkyl and (C<sub>1</sub>-C<sub>4</sub>)alkoxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, where heterocyclyl in the definition of R<sup>1</sup> to R<sup>17</sup> is a radical of a heterocyclic saturated, unsaturated or heteroaromatic ring having 3 to 6 ring atoms and 1 to 3 heteroatoms selected from the group consisting of N, O and S

where heteroaryl in the definition of R<sub>1</sub> to R<sub>17</sub> is a radical of a heteroaromatic ring having 5 to 6 ring atoms and 1 to 3 heteroatoms selected from the group consisting of N, O and S and where the substituents for substituted phenyl or substituted heteroaryl are one or more radicals selected from the group consisting of halogen, nitro, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)hydroxyalkyl and (C<sub>1</sub>-C<sub>4</sub>)alkoxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, or

R<sup>2</sup>, R<sup>3</sup> together with the carbon atom of the group R<sup>2</sup>R<sup>3</sup>C= are a non-aromatic carbocyclic ring or a heterocyclic ring having 3 to 9 ring atoms and 1 to 4 heteroring atoms selected from the group consisting of N, O and S, which ring is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, nitro, hydroxyl, oxo, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy and (C<sub>1</sub>-C<sub>4</sub>)alkylthio, or

R<sup>5</sup>, R<sup>6</sup> together with the carbon atom and the adjacent oxygen atoms of the group

C(OR<sup>5</sup>)(OR<sup>6</sup>)(OR<sup>7</sup>) are a saturated or unsaturated non-aromatic heterocyclic ring having-3 to 9 ring atoms and 1 to 4 heteroring atoms selected from the group consisting of N, O, P, and S, which ring is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, nitro, hydroxyl, oxo, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy and (C<sub>1</sub>-C<sub>4</sub>)alkylthio, or the group C(OR<sup>5</sup>)(OR<sup>6</sup>)(OR<sup>7</sup>) together is a bicyclic radical of the formula



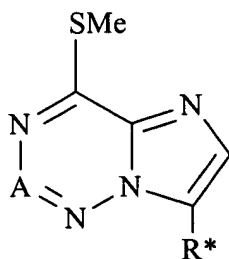
which comprises reacting a compound of the formula (IV)



with a chlorinating agent and cyclizing it to give the compound of the formula (III-1).

10. (Previously Presented) A compound of the formula (V) as set forth in claim 9.

11. (Currently Amended) A process for preparing a compound of the formula (VI)



(VI)

in which  $R^* = Z$  or G-L, Z is a precursor of the radical G-L

and A, G and L are defined further below

A is a nitrogen atom or a group of the formula C-R, where R is as defined further below,

R, is a hydrogen atom, amino, hydroxyl, mercapto, cyano, halogen, azido, nitro, SF<sub>5</sub>, unsubstituted or substituted aminosulfonyl, acyl, acylamino, acyloxy, acylthio, mono- or di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, mono- or di(C<sub>3</sub>-C<sub>9</sub>)cycloalkylamino, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>2</sub>-C<sub>4</sub>)alkenylthio, C<sub>2</sub>-C<sub>4</sub>alkynylthio, (C<sub>3</sub>-C<sub>9</sub>)cycloalkylthio, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenylthio, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfinyl, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfonyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>2</sub>-C<sub>4</sub>)alkenyloxy, (C<sub>2</sub>-C<sub>4</sub>)alkynyloxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyloxy, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>2</sub>-C<sub>4</sub>)alkenyl, (C<sub>2</sub>-C<sub>4</sub>)alkynyl, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyl, (C<sub>1</sub>-C<sub>4</sub>)alkylaminosulfonyl or di[(C<sub>1</sub>-C<sub>4</sub>)alkyl]aminosulfonyl, where each of the 23 last-mentioned radicals is unsubstituted or substituted in the hydrocarbon moiety by one or more radicals selected from the group consisting of halogen, hydroxyl, amino, nitro, formyl, carboxy, cyano, thiocyanato, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)haloalkylthio, mono(C<sub>1</sub>-C<sub>4</sub>)alkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>3</sub>-C<sub>9</sub>)cycloalkylamino, [(C<sub>1</sub>-



C<sub>4</sub>)alkyl]carbonyl, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonyl, aminocarbonyl, mono(C<sub>1</sub>-

C<sub>4</sub>)alkylaminocarbonyl and di(C<sub>1</sub>-C<sub>4</sub>)alkylaminocarbonyl,

- G is a divalent straight-chain saturated or unsaturated hydrocarbon bridge having 4  
to 6 carbon atoms in the chain, in which one or more chain members, in each case  
independently of one another, can be replaced by O, S, NH, (C<sub>1</sub>-C<sub>4</sub>)alkyl-N or  
acyl-N or, in the unsaturated case, one or more CH groups can in each case be  
replaced by a nitrogen atom, where the bridge in question is unsubstituted or  
(a) substituted by one or more identical or different radicals selected from the  
group consisting of halogen, nitro, radicals of the formula R<sup>1</sup> which are  
different from hydrogen, radicals of the formula R<sup>2</sup>R<sup>3</sup>C= and radicals of  
the formula L\*, where R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and L\* are as defined further below,  
(b) carries two or four substituents, of which in each case two together with  
the linking bridge moiety form a carbocyclic or heterocyclic ring having 3  
to 7 ring atoms, where in the case of a heterocycle the heteroatoms,  
preferably 1, 2 or 3 heteroatoms, are selected from the group consisting of  
N, O and S and where the ring in question may also have fused-on rings  
and is otherwise unsubstituted or substituted by one or more identical or  
different radicals selected from the group consisting of halogen, nitro,  
radicals of the formula R<sup>1</sup> which are different from hydrogen, radicals of  
the formula L\* and oxo, where R<sup>1</sup> and L\* are as defined further below,  
(c) is linked cyclically with L via a second direct bond or via a heteroatom  
selected from the group consisting of O and S,  
(d) has two or more substituents from the above groups (a) to (c) together,

L, L\* independently or one another are each OR<sup>4</sup>, SR<sup>4</sup>, CN, tetrazolo,

C(OR<sup>5</sup>)(OR<sup>6</sup>)OR<sup>7</sup>), -Z<sup>1</sup>, -O-Z<sup>2</sup> or -NH-Z<sup>2</sup>, where R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, Z<sup>1</sup> and Z<sup>2</sup> are as defined further below and where L may be attached cyclically to the bridge G via a second direct bond or via a heteroatom selected from the group consisting of N, O and S,

Z<sup>1</sup> is a radical of the formula COOR<sup>8</sup>, CS-OR<sup>8</sup>, CO-SR<sup>8</sup>, CS-SR<sup>8</sup>, CO-NR<sup>9</sup>-SO<sub>2</sub>-R<sup>8</sup>, CO-NR<sup>10</sup>R<sup>11</sup>, CS-NR<sup>10</sup>R<sup>11</sup>, CO-R<sup>12</sup>, CS-R<sup>12</sup>, SO-R<sup>12</sup>, SO<sub>2</sub>R<sup>12</sup>, SO<sub>3</sub>R<sup>8</sup>, SO<sub>2</sub>NR<sup>10</sup>R<sup>11</sup>, SO<sub>2</sub>NR<sup>9</sup>COR<sup>12</sup>, SO<sub>2</sub>NR<sup>9</sup>COOR<sup>12</sup>, P(=O)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=S)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=O)(R<sup>15</sup>)(OR<sup>14</sup>), P(=O)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>), P(=O)(NR<sup>10</sup>R<sup>11</sup>)-(NR<sup>16</sup>R<sup>17</sup>), P(=S)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>) or P(=S)(NR<sup>10</sup>R<sup>11</sup>)(NR<sup>16</sup>R<sup>17</sup>),

Z<sup>2</sup> is a radical of the formula COOR<sup>8</sup>, CS-OR<sup>8</sup>, CO-SR<sup>8</sup>, CS-SR<sup>8</sup>, CO-NR<sup>9</sup>-SO<sub>2</sub>-R<sup>8</sup>, CO-NR<sup>10</sup>R<sup>11</sup>, CS-NR<sup>10</sup>R<sup>11</sup>, CO-R<sup>12</sup>, CS-R<sup>12</sup>, SO-R<sup>12</sup>, SO<sub>2</sub>R<sup>12</sup>, SO<sub>3</sub>R<sup>8</sup>, SO<sub>2</sub>NR<sup>10</sup>R<sup>11</sup>, SO<sub>2</sub>NR<sup>9</sup>COR<sup>12</sup>, SO<sub>2</sub>NR<sup>9</sup>COOR<sup>12</sup>, P(=O)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=S)(OR<sup>13</sup>)(OR<sup>14</sup>), P(=O)(R<sup>15</sup>)(O<sup>14</sup>), P(=O)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>), P(=O)(R<sup>10</sup>R<sup>11</sup>)-(NR<sup>16</sup>R<sup>17</sup>), P(=S)(OR<sup>13</sup>)(NR<sup>10</sup>R<sup>11</sup>) or P(=S)(NR<sup>10</sup>R<sup>11</sup>)(NR<sup>16</sup>R<sup>17</sup>),

R<sup>1</sup> to R<sup>17</sup> independently of one another are each a hydrogen atom, (C<sub>1</sub>-C<sub>6</sub>)alkyl, (C<sub>2</sub>-C<sub>6</sub>)alkenyl, (C<sub>2</sub>-C<sub>6</sub>)alkynyl, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyl, phenyl or heterocyclyl, where each of the last-mentioned carbon-containing radicals is unsubstituted or substituted-by one or more radicals selected from the group consisting of amino, hydroxyl, mercapto, cyano, halogen, azido, nitro, SF<sub>5</sub>, aminosulfonyl, acyl, acylamino, acyloxy, acylthio, [(C<sub>1</sub>-C<sub>4</sub>)alkoxy]carbonyl, mono(C<sub>1</sub>-C<sub>4</sub>)alkylamino, mono(C<sub>3</sub>-C<sub>9</sub>)cycloalkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>2</sub>-C<sub>4</sub>)alkenylthio, (C<sub>2</sub>-C<sub>4</sub>)alkynylthio, (C<sub>3</sub>-C<sub>9</sub>)cycloalkylthio,

(C<sub>5</sub>-C<sub>9</sub>)cycloalkenylthio, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfinyl, (C<sub>1</sub>-C<sub>4</sub>)alkylsulfonyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>2</sub>-C<sub>4</sub>)alkenyloxy, (C<sub>2</sub>-C<sub>4</sub>)alkynyloxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkoxy, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyloxy, (C<sub>3</sub>-C<sub>9</sub>)cycloalkyl, (C<sub>5</sub>-C<sub>9</sub>)cycloalkenyl, phenyl, substituted phenyl, heteroaryl, substituted heteroaryl and, in the case of cyclic radicals, also by (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>2</sub>-C<sub>4</sub>)alkenyl, (C<sub>2</sub>-C<sub>4</sub>)alkynyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>2</sub>-C<sub>4</sub>)haloalkenyl, (C<sub>2</sub>-C<sub>4</sub>)haloalkynyl, (C<sub>1</sub>-C<sub>4</sub>)hydroxyalkyl and (C<sub>1</sub>-C<sub>4</sub>)alkoxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, where heterocyclyl in the definition of R<sup>1</sup> to R<sup>17</sup> is a radical of a heterocyclic saturated, unsaturated or heteroaromatic ring having 3 to 6 ring atoms and 1 to 3 heteroatoms selected from the group consisting of N, O and S

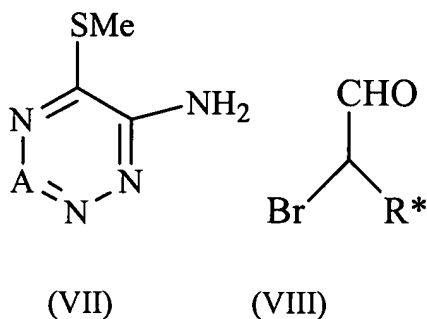
where heteroaryl in the definition of R<sup>1</sup> to R<sup>17</sup> is a radical of a heteroaromatic ring having 5 to 6 ring atoms and 1 to 3 heteroatoms selected from the group consisting of N, O and S and where the substituents for substituted phenyl or substituted heteroaryl are one or more radicals selected from the group consisting of halogen, nitro, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy, (C<sub>1</sub>-C<sub>4</sub>)alkylthio, (C<sub>1</sub>-C<sub>4</sub>)hydroxyalkyl and (C<sub>1</sub>-C<sub>4</sub>)alkoxy(C<sub>1</sub>-C<sub>4</sub>)alkyl, or

R<sup>2</sup>, R<sup>3</sup> together with the carbon atom of the group R<sup>2</sup>R<sup>3</sup>C= are a non-aromatic carbocyclic ring or a heterocyclic ring having 3 to 9 ring atoms and 1 to 4 heteroring atoms selected from the group consisting of N, O and S, which ring is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, nitro, hydroxyl, oxo, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy and (C<sub>1</sub>-C<sub>4</sub>)alkylthio, or

R<sup>5</sup>, R<sup>6</sup> together with the carbon atom and the adjacent oxygen atoms of the group

C(OR<sup>5</sup>)(OR<sup>6</sup>)(OR<sup>7</sup>) are a saturated or unsaturated non-aromatic heterocyclic ring having-3 to 9 ring atoms and 1 to 4 heteroring atoms selected from the group consisting of N, O, P, and S, which ring is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, nitro, hydroxyl, oxo, (C<sub>1</sub>-C<sub>4</sub>)alkyl, (C<sub>1</sub>-C<sub>4</sub>)haloalkyl, (C<sub>1</sub>-C<sub>4</sub>)alkoxy, (C<sub>1</sub>-C<sub>4</sub>)haloalkoxy and (C<sub>1</sub>-C<sub>4</sub>)alkylthio, or the group C(OR<sup>5</sup>)(OR<sup>6</sup>)(OR<sup>7</sup>) together is a bicyclic radical of the formula

which comprises condensing and cyclizing a compound of the formula (VII) with a compound of the formula (VIII)



12. (Previously Presented) A compound of the formula (VI) as set forth in claim 11.
  
13. (Previously Presented) A herbicidal or plant-growth-regulating composition, comprising one or more compounds of the formula (I), their salts, their tautomers or their water addition products as set forth in claim 1 and formulation auxiliaries which are customary in crop protection.

14. (Previously Presented) A method for controlling harmful plants or for regulating the growth of plants, which comprises applying an effective amount of one or more compounds of the formula (I), their salts, their tautomers or their water addition products as set forth in claim 1 onto the plants, parts of plants, plant seeds or the area under cultivation.
15. (Previously Presented) A method for controlling harmful plants and for regulating the growth of plants which comprises using a compound of the formula (I), its salt, its tautomer or its water addition product as set forth in claim 1 as herbicide or plant growth regulator.
16. (Previously Presented) The method as claimed in claim 15, wherein the compound of the formula (I), its salt, its tautomer or its water addition product is employed for controlling harmful plants or for regulating the growth in corps of useful or ornamental plants.
17. (Previously Presented) The method as claimed in claim 16, wherein the crop plants are transgenic crop plants.
18. (Cancelled)
19. (Cancelled)
20. (Cancelled)
21. (Cancelled)